Intracranial intraosseous hydatid cyst

Case report and review of the literature

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Cerebral hydatidosis accounts for approximately 1 to 3% of all cases of hydatid disease. Generally, cerebral hydatid cysts are single lesions located in the watershed of the middle cerebral artery. Primary intracranial extracerebral hydatid cysts are extremely rare. Only 2% of hydatid cysts are localized in the skeleton, and of these 3 to 4% are found in the skull.

The authors describe the case of a 10-year-old boy who was admitted to their clinic with headache and unilateral focal epileptic seizures. Computerized tomography scanning revealed a right parietal intraosseous hydatid cyst. A case of cranial intraosseous hydatid disease is presented, and the differential diagnosis and treatment are discussed in the light of literature.

KEY WORDS • echinococcosis • hydatid cyst • skull

Hydatid disease is caused by infestation of larvae of the tapeworm *Echinococcus granulosus*. Although hydatid disease has been entirely eradicated in some countries, it remains a serious health problem in certain parts of the world and is endemic in the Middle East, Mediterranean, South America, and Australia.^{1,9,12,20,21}

An adult tapeworm lives in the small intestine of definitive hosts, and its eggs contaminate water and food through ingestion of an intermediate host such as sheep, cattle, and humans.^{3,9,12} Humans are generally an incidental intermediate host, infected by the parasite frequently through the ingestion of the ova and rarely by inhalation. Hegzegan embryo becomes free after the digestion of ova in the gastrointestinal tract, and the resultant embryo most often settles in the liver through portal circulation. The embryo, which passes through the liver and crosses to the lymphatic system, enters the systemic circulation and may involve various organs, where it can cause hydatid disease.^{1,12,16,21} The most commonly affected organs are the liver and lung.

We present a case of primary cranial intraosseous hydatid cyst, and the differential diagnosis and treatment are discussed.

CASE REPORT

Presentation. This 10-year-old boy was admitted to our clinic with a history of headache and left-sided focal motor seizures. The patient recounted a history of contact with the neighbor's dog.

Examination. Neurological and physical examinations demonstrated normal results. Analysis of peripheral blood samples showed findings within the normal range, except for the presence of an erythrocyte sedimentation rate of 40 mm in the 1st hour and eosinophilia. A cystic lesion was detected in the right parietal area on x-ray films (Fig. 1). Radiological examination of the other organ systems demonstrated normal findings. Axial CT scanning of the cranium revealed a large, single intraosseous cystic lesion in the right parietal area (Fig. 2).

Operation and Treatment. After the diagnosis was made, a parietal craniectomy was performed and the intraosseous cyst was totally removed without complication (Fig. 3). The patient was discharged 7 days after the operation and received albendazole treatment.

Histopathological evaluation of the cystic lesion confirmed the diagnosis of *Echinococcus granulosus*. Albendazole treatment was initiated at a dose of 10 mg/kg three times per day. The dose was repeated twice at 4-week intervals and was stopped 3 months postoperatively. During

Abbreviation used in this paper: CT = computerized tomography.

Fig. 1. Lateral x-ray film of the skull, revealing a large cystic lesion in the right parietal area.

a 12-month follow-up period, clinical, serological, and radiological examinations demonstrated no evidence that the hydatid disease involved any other organ system.

DISCUSSION

Hydatid disease is rarely localized in the brain and comprises only 1 to 3% of reported cases.^{1,5,7-9,13,14,21} A primary cyst is the most common type of hydatid disease and the primary lesion is always solitary, but secondary disease usually involves multiple cysts that develop after embolization of a cardiac cyst, which ruptures in the left ventricle; secondary cysts are also caused by spontaneous, traumatic, or surgically induced rupture of a primary central nervous system cyst.²¹

Cerebral hydatid cysts are usually located in the distribution of the middle cerebral artery because of the embolic nature of the infestation.^{2,6,9,19,22} Because the physiological blood flow of the brain is mainly supplied by the internal carotid artery system, the migration of larvae by way of the external carotid artery system is very rare.^{3,18}

Primary intracranial extracerebral hydatid cysts have been reported but are extremely rare. They occur in three forms:^{2,3,17} 1) cranial—usually the osseous spongiosa is the first to be involved, with only 2% of the hydatid cysts localized to the skeleton and only 3–4% of these are in the skull; 2) cranial extradural—the extradural space may be infected by embolization of scolices or embryos via blood vessels,³ by extrusion of intracerebral cysts via healthy dura mater,⁴ or by erosion of osseous hydatid lesion into the extradural space;¹⁸ and 3) combined location—there may be simultaneous intracerebral, extradural, and osseous cysts.³

Computerized tomography scanning is more specific for identifying osseous lesions than other imaging modalities. In our case, cranial CT scans demonstrated the classical finding (intraosseous cystic lesion, solitary, regular contours, and spherical shape^{2,4}). In particular eosinophilic granuloma, cystic fibrous dysplasia, and calvarial epidermoid tumor/cyst must be considered in the differential diagnosis of intraosseous hydatid cyst. Eosinophilic granuloma is usually solitary and involves the diploë. Computerized tomography scanning demonstrates a

Fig. 2. Axial CT scans demonstrating intraosseous hydatid cyst in the right parietal area. Upper: Noncontrast scan. Lower: Bone window scan.

sharply margined lytic skull defect more frequently involving the outer table than the inner table.¹⁰ The cystic form of fibrous dysplasia is usually homogeneously hypodense and typically has a smooth sclerotic border. An epidermoid tumor or cyst of calvarium has a sclerotic rim and lacks inner trabeculae.¹⁰ Less likely considerations include solitary metastasis, brown tumor, and plasmacytoma. Metastases and brown tumors are rarely solitary, and they are characteristically poorly margined. Plasmacytoma may be solitary; it usually has a septated, bubbly, and irregularly margined appearance.¹⁰

Despite the use of CT scanning, the differential diagnosis of intraosseous hydatid cyst may be difficult. The Casoni intradermal test, peripheral blood and cerebrospinal fluid eosinophilia, and serological tests may be helpful for the diagnosis of hydatid disease. As in our case, in which intracranial hydatid disease was present without any other

Fig. 3. Photograph of the excised right parietal hydatid cyst.

foci or infestation, these tests may frequently yield negative findings.¹⁵ Examinations of the chest and abdominal organs with radiography and ultrasonography, respectively, must be performed routinely to exclude hydatid disease of lung and liver.

Surgical treatment is usually required when hydatid disease involves the central nervous system. Total resection of the cyst without rupture via the craniectomy is the recommended procedure.^{1,4,5,7,9,13} Extradural hydatid cysts of intraosseous origin, however, may be difficult to excise completely because of their adherence to dura and osseous trabeculae.¹¹ For this reason, if the cyst involves the extradural space, magnetic resonance imaging should be performed to obtain a diagnosis and plan the surgery. If the cyst ruptures intraoperatively, extensive irrigation of the surgical field with a hypertonic solution and a course of oral albendazole may help to prevent the recurrence.

Patients with intraosseous hydatid cysts should be treated with albendazole for at least 3 months and monitored for up to 12 months before they are considered cured.

Although primary cranial intraosseous hydatid cysts are extremely rare, hydatid disease must be considered in the differential diagnosis of intraosseous cystic lesion, especially in areas in which the disease is endemic.

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